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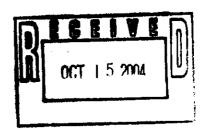
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Subject: FINAL REPORT FOR THE SOIL SAMPLES FROM 216-S-20 CRIB ( 222S20040166

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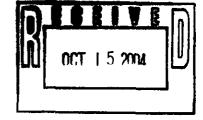
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October 13, 2004

CH2M-0403032

Mr. S. J. Trent, Manager Environmental Information Systems Fluor Hanford, Inc. Post Office Box 1000 Richland, Washington 99352-0450



Dear Mr. Trent:

#### FINAL REPORT FOR THE SOIL SAMPLES FROM 216-S-20 CRIB-222S20040166

- References:
- M. E. Todd-Robertson et al., "Analytical Instruction: 200-LW-1 and 200-LW-2 Operable Unit Characterization Sampling and Analysis Concurrence for Analytical Requirements," Fluor Hanford Groundwater Remediation Project, dated September 1, 2004.
- 2. HNF-SD-CP-QAPP-016, 222-S Laboratory Quality Assurance Plan, Revision 8, dated January 29, 2004.
- 3. Letter, H. L. Anastos, FH, to Distribution, "Semi-Volatile Organic Compound Analysis," FH-0300526, dated February 3, 2003.
- 4. Letter, H. L. Anastos, FH, to Distribution, "Volatile Organic Compound Analysis," FH-0300583, dated February 3, 2003.
- RPP-6268, TWINS/Labcore Configuration Control Desk Manual, Revision 1, CH2M HILL Hanford Group, Inc., Richland, Washington, dated April 19, 2002.
- 6. SW-846, Test Methods for Evaluating Solid Waste: Physical/Chemical Methods, Third Edition, as amended, U.S. Environmental Protection Agency, Washington, D.C., September 1986.

This letter and four attachments represent the final analytical data report for the soil samples from the 216-S20 crib, which were received at the 222-S Laboratory on August 30, 2004. The samples were analyzed in accordance with Reference 1 through Reference 5.

Should you have any questions regarding this matter, please contact me at 373-4314.

Very truly yours,

R. A. Bushaw, Project Coordinator Analytical Project Management

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Attachments (4)

Attachment 1

NARRATIVE

Consisting of 7 pages, including coversheet

# 222-S LABORATORY FINAL REPORT FOR THE SOIL SAMPLES FROM 216-S-20 CRIB-222S20040166

#### 1.0 INTRODUCTION

Two soil samples from the 216-S-20 crib were received at the 222-S Laboratory on August 30, 2004. The samples were analyzed in accordance with the 200-LW-1 and 200-LW-2 Operable Unit Characterization Sampling and Analysis Concurrence for Analytical Requirements (analytical instructions) (Reference 1), 222-S Laboratory Quality Assurance Plan (QAPP-016, Reference 2), "Semi-Volatile Organic Compound Analysis" (Reference 3), and "Volatile Organic Compound Analysis" (Reference 4), referenced in the cover letter.

A Data Summary Report is included as Attachment 2. The correlation between the customer sample identification number and laboratory identification numbers is presented in the Sample Breakdown Diagrams included as Attachment 3. Copies of the receipt paperwork are included as Attachment 4.

#### 2.0 SAMPLE APPEARANCE AND HANDLING

Sample delivery group 222S20040166 consists of customer samples B191F1 and B191J2. The samples were collected on August 18, 2004, but were not delivered to the laboratory until August 30, 2004. The samples were described as dark brown, medium coarse sand with some small stones.

Except for analysis for volatile organic compounds (VOC), the samples were stirred with a spatula prior to removing aliquots for analysis. With this type of sample, this method is typically not sufficient to achieve homogenization. Typically, the effectiveness of the homogenization can be determined by evaluating the relative percent difference (RPD) between sample and duplicate results. However, since many of the requested analytes were not detected in the sample, an RPD wasn't calculated for all analytes. A further discussion of homogeneity is included in Section 5.3.

For the VOC analysis, three 40-mL bottles were filled to the top with soil, leaving no head space. No preservative was added to the sample bottles in the field. With the sample received in this configuration, the bottles had to be opened in a hood to obtain aliquots for analysis. To reduce the time that the sample was exposed to the atmosphere, it was not stirred prior to removing the aliquots. However, opening the sample to the atmosphere for even a short period of time compromises the integrity of the sample and the results may be biased low.

#### 3.0 HOLDING TIMES

The analytical instructions requested that the laboratory make every effort to meet analytical holding times (SW-846, Reference 6 in the cover letter) for all analyses. The holding times were met for all analyses except for pH, sulfide, and cyanide (CN). For pH (24 hours) and sulfide (7 days), the holding times were not met because these samples were not delivered to the laboratory until 12 days after sampling. Reagents for the CN analysis (14-day holding time) have a short shelf life and, therefore, were ordered only after the samples were received (there was no advance notice for these samples). Since the samples were received on day 12 of the 14-day holding time, there was insufficient time to order and prepare the reagents and begin the analysis within the holding time.

#### 4.0 ANALYTICAL RESULTS

The Data Summary Report in Attachment 2 presents the analytical results for the requested analytes. In this table, solid samples that were prepared by water digest are indicated with a "W" in the aliquot class (A#) column, and an "S" indicates a distillation preparation was used. If there is no letter identifier in this column, it indicates that the analysis was performed on a direct subsample with no separate preparation or with sample preparation that was included as part of the analytical procedure steps.

Note that the "Unit" column in the Data Summary Report indicates the units for the sample results. The reporting units for the blank do not all match those for the sample results. The units for the blank are  $\mu$ g/mL for the anions and ammonium reported by ion chromatography (IC), as well as for CN, sulfide, and mercury.

#### 5.0 QUALITY CONTROL RESULTS

#### 5.1 LABORATORY CONTROL SAMPLES

The accuracy of the analysis was evaluated from the recovery of a laboratory control sample (LCS) and a matrix spike (MS). The requested accuracy was 70-130% recovery. The requested accuracy of the LCS for the pH analysis was ±0.1 pH units.

All LCS recoveries were acceptable in accordance with the analytical instructions and QAPP-016. The LCS for the pH analysis also met the requested criterion.

For the semivolatile organic compound (SVOC) and the VOC analyses, the analytical instructions only required the set of compounds indicated in the letter from H. L. Anastos (Reference 3). However, tributyl phosphate was added to the "typical" standard mix used for LCS for the SVOC analysis because it was a requested analyte for reporting. Since this compound is not on the "typical" reporting list, the quality control (QC) parameters for this compound were not set up in the laboratory database, and the QC results for the LCS are not included in the Data Summary Report. The LCS recovery was 122% for the tributyl phosphate.

For the VOC analysis, a ketone mix (containing 4-methyl-2-pentanone, 2-butanone, and acetone) and butanol were part of the standard mix used for the LCS in addition to the "typical" set of compounds indicated in the letter from H. L. Anastos (Reference 4). These compounds are part of the QC protocol associated with an unrelated project. Although the LCS and MS recoveries for ketones and 1-butanol were not required to be reported, they are included in the Data Summary Report because they are compounds requested in the analytical instructions.

#### 5.2 METHOD AND PREPARATION BLANKS

For most analyses, no analytes were detected in the method or preparation blank. However, F and Cl were detected in the blank that was prepared and analyzed with the IC anion analysis. The level of Cl detected in the sample was the same as that detected in the preparation blank. The level of F detected in the sample was only about 3.5 times higher than that detected in the preparation blank. These levels of contamination are considered significant and the results should be considered biased high due to the contamination level. The sample was not reprepared or reanalyzed because the reported results were lower than the level of concern for this project and concurrence was received from the customer point of contact to report the biased results.

#### 5.3 DUPLICATE ANALYSES

One duplicate sample was analyzed for each method. The requested precision for analysis was  $\pm 30\%$  for all methods except pH. For pH, a precision of  $\pm 0.1$  pH units was requested. The precision was assessed by calculating the RPD between the sample and duplicate results. For VOC, SVOC, and the polychlorinated biphenyl (PCB) analyses, the analysis precision was evaluated by calculating the RPD between an MS and a matrix spike duplicate (MSD). An RPD <30% meets the requested precision. Although an RPD was calculated and reported for the pH analysis, the results were also evaluated against the requested criterion of  $\pm 0.1$  pH units. The analysis met the requested precision.

All analyte results met the RPD criterion stated in the analytical instruction. However, note that for phosphate and sulfate, the sample portion was reported as less than the detection limit and the duplicate had positive results. Therefore, as discussed previously, no RPD was calculated for these analytes. For the phosphate, the duplicate result was only slightly above the detection limit. However, for sulfate, the sample result was <12.8  $\mu$ g/g and the duplicate had a positive result of 21.2  $\mu$ g/g. The duplicate result is less than two times the detection limit and should be considered an estimate due to the reduced precision of the analysis near the detection limit. The large difference between the sample and duplicate results is likely a consequence of the inhomogeneity of the sample. The laboratory does not have appropriate equipment to effectively homogenize a sample that contains both sand and stones, so no reanalysis was requested.

The Data Summary Report does not report the RPDs for the MS/MSD analysis. This information is provided in Table 1 for VOC, Table 2 for SVOC, and Table 3 for PCB analysis.

Table 1. MS/MSD Recoveries and RPD for VOC.

Compound	MS (%)	MSD (%)	RPD (%)
Benzene	110	110	0
Chlorobenzene	110	109	1
1,1-Dichloroethene	98	97	1
Toluene	112	111	1
Trichloroethene	94	95	1
Acetone	80	85	6
2-Butanone	78	83	6
4-Methyl-2-pentanone	110	114	4
1-Butanol	62	73	16

Table 2. MS/MSD Recoveries and RPD for SVOC.

Compound	MS (%)	MSD (%)	RPD (%)
Phenol	81	74	9
2-Chlorophenol	78	70	11
1,4-Dichlorobenzene	82	70	16
N-Nitroso-di-n-propylamine	80	70	13
1,2,4-Trichlorobenzene	82	72	13
4-Chloro-3-methylphenol	88	84	5
Acenaphthene	89	81	9
4-Nitrophenol	100	96	4
2,4-Dinitrotoluene	97	92	5
Pentachlorophenol	100	95	5
Pyrene	108	100	8
Tributyl phosphate	111	119	7

Table 3. MS/MSD Recoveries and RPD for PCB.

Compound	MS (%)	MSD (%)	RPD (%)
Aroclor-1254	108	124	14

#### 5.4 MATRIX SPIKE AND MATRIX SPIKE DUPLICATE

Where applicable, one MS sample was analyzed for each method. An MS analysis was not applicable for the pH analysis. For PCB analysis, only Aroclor-1254 is included in the MS because it is the aroclor most commonly detected in samples on the Hanford Site. For the SVOC analysis, tributyl phosphate was not listed in the letter from H. L. Anastos (Reference 3) but was added to the standard mix for the MS. As discussed for the LCS, the MS recovery also was not included in the Data Summary Report, but is included in Table 2. For the VOC analysis, the MS contained the same "non-typical" compounds as discussed for the LCS, and the recoveries are included in the Data Summary Report and in Table 1.

As stated in Section 5.1, the accuracy of the analyses was evaluated from the LCS and MS recoveries, with a requested accuracy of 70-130% recovery. All MS and MSD recoveries met the criterion listed in the analytical instructions, except the MS for 1-butanol. The MS recovery for 1-butanol was low (62%) and the recovery for the MSD (73%) was just inside the requested limits. The sample, MS, and MSD were analyzed twice, with low recoveries each time. Therefore, the low recovery was attributed to matrix interference, so no further reanalysis was requested. The matrix interference could cause a low bias in the quantitation for the 1-butanol.

The Data Summary Report does not report the recoveries for the MSD analyses. This information is provided in Tables 1, 2, and 3.

#### 5.5 SURROGATE RECOVERIES

Surrogate standards are added to all field and QC samples for VOC, SVOC, and PCB analyses. The surrogate is added to monitor total method recovery through preparation, sample matrix cleanup, and analysis. The surrogate recovery for the preparation blank analyzed for PCB was high, which would indicate a high bias. However, since no PCBs were detected in the blank and the surrogate recoveries for the sample, MS, and MSD all met the acceptance criteria, the results should be considered accurate as reported. One of the six surrogates analyzed with the SVOC (2,4,6-tribromophenol) had slightly high recoveries for all field and QC samples. The recoveries were outside of the statistical limits but within the administrative limits for the method. No reanalysis was requested because the MS and MSD recoveries were all within the acceptance limits, indicating that the accuracy of the analysis was acceptable. The high surrogate recoveries did not affect the usability of the reported results. All surrogate recoveries for the VOC analysis met the requirements in QAPP-016.

#### 5.6 TENTATIVELY IDENTIFIED COMPOUNDS

The analytical instruction lists n-butyl benzene as a target compound for VOC analysis. The laboratory does not routinely report results for this compound, as indicated in the letter from H. L. Anastos (Reference 4). Review of the data indicated that n-butyl benzene was not detected in the sample. Also, no other tentatively identified compounds were detected.

#### 5.7 TARGET QUANTITATION LIMITS

The analytical instructions listed target quantitation limits (TQL) for each requested analyte. The Data Summary Report provides method detection limits (MDL). These must be converted to estimated quantitation limits (EQL) to compare with the requested TQLs. The EQL is calculated as 10 times the reported MDL. Analytes that are detected above the MDL but are less than the EQL are qualified with a "J" flag, as specified in the TWINS/Labcore Configuration Control Desk Manual (Reference 5 in the cover letter), to indicate that the result should be considered an estimate because the concentration is less than that which can be reliably achieved within the specified limits of precision and accuracy.

The laboratory EQLs did not all meet the requested TQLs because the procedures and equipment have been altered to handle small sample sizes to reduce the level of radioactive exposure to the analyst. For the PCB and Hg analyses, the reported concentrations are greater than 10 times the laboratory MDL and the requested TQL. Therefore, these results should be considered accurate, regardless of the EQL. For the other analyses, the laboratory used the largest feasible sample sizes to obtain the lowest detection limits possible.

#### 6.0 ANALYTICAL PROCEDURES

Table 4 presents the 222-S Laboratory analytical procedures used to generate the reported results.

Table 4. Analytical Procedures.

	Preparation Erocciano	Analysis Precodure
	Inorganic Analyses	
pН	Direct	LA-212-105 Rev. D-0
Hg	Direct	LA-325-106 Rev. C-1
CN	Direct	LA-695-102 Rev. I-2
NH <sub>4</sub>	Distillation	LA-533-101 Rev. L-0
IC	Water digest	LA-533-107 Rev. D-0
Sulfide	Direct	LA-361-101 Rev. A-2
	Organic Analyses	
VOC	Direct	LA-523-118 Rev. A-2
SVOC	Organic extraction	LA-523-135 Rev. A-2
PCB	Organic extraction	LA-523-140 Rev. C-0

Notes:

Water digest procedure: LA-504-101 Rev. I-0 Distillation procedure: LA-544-112 Rev. A-1 Organic extraction procedure: LA-523-138 Rev. D-0

## Attachment 2

DATA SUMMARY REPORT

Consisting of 4 pages, including coversheet

#### Attachment 2 \$20 CRIB Data Summary Report

CORE NUMBER: 222520040166

SEGMENT #: B191F1

SEGMENT PORTION: NH4 Distillation

	1				·						
Sample# R A# Analyte_	Unit	Standard %		Result	Duplicate	Average	RPD %	Spk Rec %	Det Limit	Count Err%	Qual Flags
S04M000279 S Ammonium Ion-IC-Dionex 100	Ug/g	99.2	<0.100	<46.3	<45.4	n/a	n/a	101	46	n/a	U

SEGMENT PORTION: PCB

{·	$\Box$									-	[ · · ·		- '
Sample# R	<b>A#</b>	Analyte	Unit	Standard %	Blank	Result	Duplicate	Average	RPD X	Spk Rec %	Det Limit	Count Err%	Qual Flags
S04M000277		Aroclor-1016WET by SW-846 8082	ug/Kg	n/a	<82	<40	n/a	n/a	n/a	n/a	4.e+01	n/a	U
S04M000277		Aroclor-1221WET by SW-846 8082	ug/Kg	n/a	<26	<13	n/a	n/a	n/a	n/a	1.e+01	n/a	C
S04M000277		Aroclor-1232WET by SW-846 8082	ug/Kg	n/a	<4.6e+02	<2.3e+02	n/a	n/a	n/a	n/a	2.e+02	n/a	Ü
\$04M000277		Aroclor-1242WET by SW-846 8082	ug/Kg	n/a	<84	41	n/a	n/a	n/a	n/a	4.e+01	n/a	_
S04M000277		Aroclor-1248WET by SW-846 8082	ug/Kg	n/a	<27	<13	n/a	n/a	n/a			n/a	U
S04M000277		Aroctor-1254WET by SW-846_8082	ug/Kg	93	<16	1.7e+02	n/a	n/a	n/a	1.1e+02		n/a	
S04H000277		Aroclor-1260WET by SW-846 8082	ug/Kg	n/a	<1.1e+02	<55	n/a	n/a	n/a	n/a	5.e+01	n/a	U

SEGMENT PORTION: Parent

~-	Chiatras I dati i A	77 C . I	WI 0116												
	Sample#	R A	# Analyte	Unit	Standard X	Blank	Result	Duplicate	Average	RPD %	Spk Rec %	Det Limit	Count Err%	Qual Flags	
	S04M000274		Cyanide Water Distil	lation ug/g	107	<0.0550	<0.278	<0.275	n/a	n/a	108	0.28	n/a	U	
	S04M000274	7	Mercury by CVAA (PE)	with FIAS ug/g	103	<1.00e-04	66.6	79.8	73.2						
	S04M000274		pH on Solid Samples	DH	n/a	n/a	8.14	8.18	8.16	0.490	⊓/a	0.010	n/a		

SEGMENT PORTION: SVOA

	T				T					<u> </u>			
Sample#R	A#	Analyte	Uni t	Standard %	Blank	Result	Duplicate	Average	RPD %	Spk Rec X	Det Limit	Count Err%	Qual Flags
S04M000276	T	Pentachlorophenol	ug/Kg	93	<1.0e+03	<9.2e+02	n/a	n/a	n/a	1.0e+02	9.e+02	n/a	U
S04H000276	П	Phenol	ug/Kg	77	<1.0e+03	<9.2e+02	n/a	n/a	n/a	81	9.e+02	n/a	U
S04M000276		2-Chlorophenol	ug/Kg	74.	<1.0e+03	<9.2e+02	n/a	n/a	n/a	78	9.e+02	n/a	U
S04M000276	T	Pyrene	ug/Kg	1.0e+02	<1.0e+03	<9.2e+02	n/a	n/a	n/a	1.1e+02	9.e+02	n/a	U
S04M000276	Т	N-Nitroso-di-n-propylamine	ug/Kg	75	<1.0e+03	<9.2e+02	n/a	n/a	n/a	80			U
S04M000276	Т	1,2,4-Trichlorobenzene SV	ug/Kg	79	<1.0e+03	<9.2e+02	n/a	n/a	n/a			n/a	U
S04N000276	1	4-Chloro-3-methylphenol	ug/Kg	85	<1.0e+03	<9.2e+02	n/a	n/a	n/a	88	9.e+02	n/a	Ü
S04H000276	П	Acenaphthene	ug/Kg		<1.0e+03	<9.2e+02	n/a	n/a	n/a			n/a	U
S04M000276	Т	4-Nitrophenol	ug/Kg	96	<1.0e+03	<9.2e+02	n/a	n/a	n/a	1.0e+02	9.e+02	n/a	U
S04M000276	Т	2,4-Dinitrotoluene	ug/Kg	95	<1.0e+03	<9.2e+02	n/a	n/a	n/a			n/a	U
\$04H000276	Т	1,4-Dichlorobenzene	ug/Kg	77	<1.0e+03	<9.2e+02	n/a	n/a	n/a	82	9.e+02	n/a	U
S04M000276		Tri-n-butylphosphate	ug/Kg	n/a	<1.0e+03	<9.2e+02	n/a	n/a	n/a	n/a	9.e+02	n/a	U

04M0dD273	A# Analyte	Unit	Standard %	Blank		Duplicate	Average				Count Err%	Qual
	Methylene Chloride	ug/Kg	n/a	<1.3	<1.3		n/a	n/a	n/a		n/a	
04MQB0273	Acetone	ug/Kg	1.1e+02	<0.92	<0.92		n/a	n/a				
04M <b>#0</b> 0273	1-Butanol	ug/Kg	93	<4.0	<3.9		n/a	n/a	62		n/a	
04M000273	1.1-Dichloroethane	ug/Kg	n/a	<0.80	<0.80		n/a			0.8		
04MQQQ273	1,2-Dichloroethene (cis	2 tron un/Ka	n/a	<1.4	<1.4	n/a	n/al	n/a	n/a	1	n/a	



30-sep-2004 10:53:23 A-0002-1(23)

Sample#	R A#	Analyte	Unit	Standard %	Blank	Result	Duplicate	Average	RPD %	Spik Rec %	Det L <u>i</u> mit	Count Err%	Qual Flags
S04M000273		Chloroform	ug/Kg	n/a	<0.72	<0.72	n/a						
S04M000273		1,2-Dichloroethane	Ug/Kg	n/a	<0.76	<0.76	n/a	n/a	n/a			n/a	U
S04M000273		2-Butanone	ug/Kg	1.0e+02	<0.82	<0.82	n/a	n/a	n/a	78	0.8	n/a	U
\$04H000273		1,1,1-Trichloroethane	ug/Kg	n/a	<0.70	<0.70	n/a	n/a	n/a	n/a	0.7	n/a	U
S04M000273		Carbon Tetrachloride	ug/Kg	_n/a	<1.3	<1.3	n/a	n/a	n/a			n/a	Ü
SO4M000273	Τ.	Trichloroethene	Ug/Kg	96	<0.86	<0.86	n/a	n/a	n/a	95	0.9	n/a	U
S04M000273		1,1,2-Trichloroethane	ug/Kg	n/a	<0.68	<0.68	n/a	n/a	n/a	n/a	0.7	n/a	כ
S04M000273		Benzene	ug/Kg	1.1e+02	<0.66	<0.66	n/a	n/a	/ <b>•</b>	1.1e+02	0.7	n/a	U
S04M000273	T	4-Nethyl-2-pentanone	ug/Kg	1.2e+02	<0.74	<0.74	n/a	n/a	r/a	1.1e+02	0.7	n/a	U
S04M000273	7	Tetrachloroethene	ug/Kg	n/a	<0.70	<0.70	n/a	n/a	n/a	n/a	0.7	n/a	U
S04M000273	7	Toluene	∪g/Kg	1.1e+02	<0.64	<0.64	n/a	n/a	_/a	1.1e+02	0.6	n/a	U
S04M000273		Chlorobenzene	ug/Kg	1.1e+02	<0.76	<0.76	n/a	n/a	r/a	1.1e+02	0.8	n/a	5
S04H000273	1	Ethylbenzene	Ug/Kg	n/a	<0.98	<0.98	n/a	n/a	n/a	n/a	1	n/a	Ū
S04H000273		Xylenes (total)	ug/Kg	n/a		<1.6	n/a	n/a	n/a	n/a	2	r/a	U
S04M000273		1.1-Dichloroethene	ug/Kg	99		<0.76			n/a	98	0.8	n/a	U

SECME	MT I	7000	ON -	Ustar	Digest	
3E UM 6				BOLE:	DINESL	

Scurent Pokiton, nater Digest													
Sample#	R A# A	inalyte	Unit	Standard X	Blank	Result	Duplicate	Average	RPD %	Spk Rec %	De <u>t Li</u> mit	Count Err% Que	al Flags
S04M000278	U F	luoride IC SW846	U9/9	96.2	0.0200	6.51	6.74	6.63			1.1	n/a <i>B</i> (	- 4
S04M000278	W C	hloride SW-846	Ug/g	98.5	0.0200	1.86	1.92	1.89	3.39		1.6		2/1/2/20
S04M000278	W N	itrite IC SW846	UQ/g	93.6	<0.108	<10.0	<10.4	n/a			10	n/a	U 3 /2 /00
S04M000278	U N	itrate by IC SW846	Ug/g	97.9	<0.139	18.6	17.3	18.0	7.15			n/a <i>B</i>	102
S04M000278	W PI	hosphate by IC \$W846	ug/g	95.4	<0.120	<11.2	11.5	n/a	n/a	95.5		n/a	3/2/05
S04M000278	W S	ulfate by IC_SW846	ug/g	94.5	<0.138	<12.8	21.2	n/a	n/a	96.9	13	n/a	U



#### Attachment 2 \$20 CRIB Data Summary Report

CORE NUMBER: 222520040166

SEGMENT #: B191J2

SEGMENT PORTION: Parent

SEGNENT FORT	Town Parent			r		T	1						٦
Sampoles# R	A# Analyte	Unit s	Standard X	Blank	Result	Duplicate	Average	RPD %	Spk Rec X	Det Limit	Count Err%	Qual Flag	8
S04MD00275	Sulfide by Microdist. & ISE	U9/9	97.2	<0.158	23.9	28.0	25.9	15.6	92.0	15	n/a	B -	₫

3/2/05

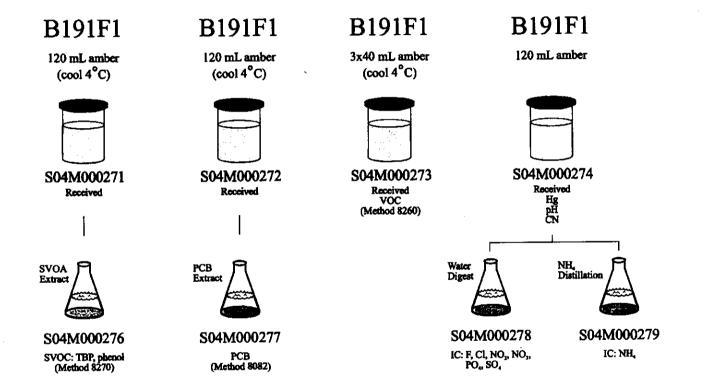


Attachment 3

SAMPLE BREAKDOWN DIAGRAM

Consisting of 2 pages, including coversheet

## S20 CRIB 216-S-20 Borehole Samples Group 222S20040166



250 ml amber (cool 4°C)



S04M000275
Received
Sulfide

## Attachment 4

## RECEIPT PAPERWORK

Consisting of 5 pages, including coversheet

	FLUC	OR Hanford Inc.		CHAIN C	F CUSTODY	/SAMPLE AP	ialysis r	<b>EQUEST</b>		F03-025-094	•	PAGE 1 OF 2				
COLLECTOR	-		COMPANY CON	TACT	TEL	EPHONE NO.		PROJECT	COORDINATOR	PRICE CODE	8N	DATA				
Pc-pe/Pfister/W			TRENT, STEVE		37	3-5689		TRENT, SJ	PRICE COPE	OIA	TURNAROUND					
SAMPLING LO	CATION.	AG- 8110/04	PROJECT DESIGNATION						SAF NO. AIR QU			45 Days / 45 Days				
		29.5' -32 '	<del></del>	Characterization -				F03-025				43 Days				
ICE CHEST NO	<b>).</b>		FIELD LOGBOOI	( NO.		COA			OF SHIPMENT							
			HNF-N-356 1			119143ES10		Governme								
SHIPPED TO		8/30/04	OFFSITE PROPE	RTY NO.				i i	ADING/AIR BILL	NO.						
Weste Samplin	i <del>g it Chara</del> c	terization 222-S	N/A		<b>-</b>			N/A								
MATRIX* A=Air DL=Drum	POSSIE N/A	LE SAMPLE HAZARDS/ REMARKS	PRESER	VATION	Cool 4C	Cool 4C	Cool 4C	Cool 4C	Mone None	. None						
Liquids DS=Drum Solids			TYPE OF O	ONTAINER	Gs*	aG	₽G ⟨	√ Gs* /	P / aG	aG aG						
L=Liquid O=Oil S=Soil			NO. OF CONTAINER(S)  VOLUME		3	1	1 /	3 soming put	1 (1) 1 (1) 250mL (2) 250mL	1 120mL						
SE=Sediment T=Tissue V=Vegitation W=Water					40mL	120mL	120mL		500mL (2) 250m	120mL						
WI-Wipe X=Other	SPECIAL HANDLING AND/OR STORAGE N/A		SAMPLE ANALYSIS		SEE ITEM (1) IN SPECIAL INSTRUCTIONS	SEE ITEM (2) IN SPECIAL INSTRUCTIONS	PCBs /CG2;	SEE IDEN (3) PH SPECIAL DISTRUCTIONS	SEE (19M (4) IN SEE SPECIAL SPECIAL INSTRUCTIONS INSTR	TEM (S) IN SEE ITEM (6) IN SPECIAL INSTRUCTIONS						
							1	1/			į					
SAMPL	E NO.	MATRIX*	SAMPLE DATE	ļ <u>.                                    </u>						,						
B191F1		SOIL	8-18-04	0858	<u> </u>	<u> </u>	1×	·×	X	<u> </u>						
		·		ļ	<u> </u>	<u> </u>			<del> </del>		ļ <u>.</u>					
						<u> </u>	ļ. <u></u>									
					<u> </u>		<u> </u>									
										<u> </u>						
CHAIN OF PO			SIGN/ PRINT	NAMES				PECIAL INST								
RELINQUISHED	BY/REMO	VED FROM DATE/TIME	RECEIVED	STORED IN		DATE/1	DME	SEE PAGE 2 FO	OR ALL SPECIAL	INSTRUCTIONS						
- 17 T/		APPERATE FIRST POR		fat	8-1	<del></del>	••									
RELINQUISHED	- 2		RECEIVED BY	STORED IN	chil	DATE/T										
KELINGHISHED			JUNEA/ART AVIII													
50/2/	7 4 KM	8-26-5	1 Christy Phinasper 8-30-64													
RELINQUISHES	BW REMO	VED FROM DATE/TIME	RECEIVED BYT/STORED IN DATE/TIME					•								
RELINQUISHED	BY/REMO	VED FROM DATE/TIME	RECEIVED BYT/STORED IN DATE/TIME				TME									
RELINQUISHED BY/REMOVED FROM DATE/TIME			RECEIVED BYT/STORED IN DATE/TIME					_				,				
RELINQUISHED BY/REMOVED FROM DATE/TIME			RECEIVED BYT/STORED IN DATE/TIME													
LABORATOR SECTION	KT j	CELVED BY					1	TITLE				DATE/TIME				
FINAL SAMPLE DISPOSAL METHOD 000001 8						DISPOSED BY				DATE/TIME						

FLUOR Hanford Inc.	CHAT	CHAIN OF CUSTODY/SAMPLE ANALYSIS REQUEST							
COL/ ECTOR Pope/Pfister/Wiberg/Tyra	COMPANY CONTACT TRENT, STEVE	TELEPHONE NO. 373-5689	PROJECT COORDINATOR TRENT, SJ	PRICE CODE 8N	DATA TURNAROUND				
SAMPLING LOCATION 216-5-20; 02305-5-5-7 24.5-32.	PROJECT DESIGNATION 200-LW-1/LW-2 Characterization	n - Soil	SAF NO. AIR QUALITY D						
ICE CHEST NO. 8 70-44	FIELD LOGBOOK NO. HNF-N-356 1	COA 119143E510	METHOD OF SHIPMENT Government Vehicle						
SHIPPED TO Waste Sampling & Characterization	OFFSITE PROPERTY NO.		BILL OF LADING/AIR BILL NO. N/A						
SPECIAL INSTRUCTIONS									
The lab is to analyze pH within 24 hours of sample rec 300.0 will not be met.	elpt. The lab is to report kerosene range orga	unics from the WTPH-D analysis. FH a		time for Nitrate, Nitrite and Pho	osphate by EPA Method				

(1)VOA - 8260A (TCL); VOA - 8260A (Add-On) {1-Butanol} (2)Semi-VOA - 8270A (TCL) {Phenol} Semi-VOA - 8270A (Add-On) {Tributyl phosphate} TPH-Diesel Range - WTPH-D (Total petroloum hydrocarbons - diesel range, Total petroloum hydrocarbons - kerosene range} TPH-Gastiline

Range-WTH'3;
(3)Alevids, Glycole, & Katanes -8015 (Cthylene glycol) P 8/34/9 1
(4)Gamma Spect-Add-on (Antimony 125, Cestum-134) Isotopic Plannium; Nephinium 237; Americium 241;
(5)ICP/MS - 200.8 (TAL) (Antimony, Barlum, Cadmium, Chromium, Copper, Mickel, Silver) ICP/MS - 200.8 (Add-on) (Arsenic, Beryllitim; Lead, Marcury, Selenium, Uranium) ICP Metals - 5010A (Add-on) (Bismuth) (BOXOY)
(6)IC Ankons - 300.0 (Chloride, Fluoride, Nitrogen in Nitrate, Nitrogen in Nitrate, Phosphate, Sulfate) Cations (IC) - 300.7 (Nitrogen in ammonium) Cyanide (Total) - 335.2; pH (Soil) - 9045; #6 B1 ate-fluid 94.71

<u> 100000009</u>

	FLUOR	Hanford Inc.		CHAIN	OF CUSTO	Y/SAMPLE AI	NALYSIS R	EQUEST		F03-025-114		PAGE 1 OF 1			
COLLECTOR Pope/Pfister/W	/iberg/Tyra		COMPANY CONTRENT, STEVE	TACT	-	LEPHONE NO. 373-5689		PROJECT (	PRICE CODE	8N		ATA IAROUND			
SAMPLING LO	CATION	180- 3-10-14	PROJECT DESIG	NATION			<del></del>	SAF NO. AIR QU					Days / Days		
216-5-20; <del>22.</del> 6	<del>n-31.51</del> Z	551-321	200-LW-1/LW-2	Characterization -	<b>Soil</b>			F03-025							
ICE CHEST NO	),		FIELD LOGBOOK NO. COA						OF SHIPMENT						
			HNF-N-356 1 119143ES10					Federal Ex	press						
HIPPED ID	<u> </u>	222-5	OFFSITE PROPE	ERTY NO.				BILL OF LADING/AIR BILL MO.							
MATRIX* A=Air XL=Drum	POSSIBLE N/A	E SAMPLÉ HÁZARDS/ REMARKS	PRESER	NOTTAN	Cool 4C	None	/								
iquids IS=Drum iolids			TYPE OF O	ONTAINER	aG	*c 3/					,				
L=Liquid D=Oil S=Soil SE=Sediment I'=Tissue V=Vegitation W=Water	SPECIAL HANDLING AND/OR STORAGE		VOLUME		1 250mL	tal									
w=water WI=Wipe X=Other					SEE ITEM (1) SPECIAL INSTRUCTION	EN SEE UEM (2) IN SPECIAL INSTRUCTIONS	!								
SAMPI	E NO.	MATRIX*	SAMPLE DATE	SAMPLE TIME											
3191J2		SOIL	8-18-54	0858	人	_1									
											<u> </u>				
	·										1				
					1										
				ļ			<del>                                     </del>								
HAIN OF PO	SSESSION		SIGN/ PRIN	T NAMES	_1	<u></u>	S	PECIAL INST	RUCTIONS	82-8/Min4	1		2-8/10/0		
RELINOUISHED RELINOUISHED	SA/c	S (F-SY ()00	RECEIVED BY	STORED IN		DATE/1 18-01 /2 DATE/1 26/64 /4:	TIME I	2)Nickel-63: (	<del>lex - 7196; NO2/I</del> <del>Samma Spec - Rac</del>	N <del>O3 - 353.2,</del> Sulfic dium (Radium 226 Tritium - H3, Car	, Radium	0; <del>Oil &amp; G</del> reas 1-228) Techne	<del>e 413.1,</del> tium 99;		
RELINQUISHER	BY/REHOVE	ED FROM DATE/TIME	RECEIVED BYT		moc	DATE	35	,							
RELINQUISHED BY/REMOVED FROM DATE/TIME			RECEIVED SYT/STORED IN			DATE/TIME									
RELINQUISHED BY/REMOVED FROM DATE/TIME			RECEIVED BYT/STORED IN			DATE/TIME						•			
RELINQUISHED BY/REMOVED FROM DATE/TIME			RECEIVED BYT/STORED IN DATE/TIME			TIME				·					
LABORATO SECTION	KT j	ETAED &A					1	TYLE				DATE/TIME			
FINAL SAM	rut	POSAL METHOD						DISPOSED BY				DATE/TIME			
			020			· · · · · · · · · · · · · · · · · · ·									

GENERATOR KNOWLEDGE INFORMATION											
1. Chain of Custody Number CACN/COA 119142/ES20 Customer Identification	fication Num	ber SAF	F03-020								
2. List generator knowledge or description of process that produced sample. Or list description of sample source			<del></del>								
Characterization samples collected from a characterization borehole drilled near the 216-S-20 waste site											
MSDS Available? No X Yes Hanford MSDS No.											
3. List all waste codes and constituents associated with the waste or media that was sampled, regardless of CEF	CLA status.		,								
a) Does the sample contain any of the following listed waste codes? By checking "unknown" the customer understands that no knowledge is available following a care.	hii seemh										
List Federal Waste Code(s): List Constituent(s):											
P Codes:	Yes	⊠ No	Unknown								
U Codes:	Yes	⊠ No	Unknown								
K Codes:	Yes	⊠ No	Unknown								
F Codes:	Yes	⊠ No	Unknown								
b) List applicable characteristic waste codes, flash point, pH, constituents, and concentrations as appropriate											
D001: ☐ FP <100°F ☐ FP ≥100 <140°F ☐ DOT Oxidizer	☐ Yes	⊠ No	Unknown								
D002: ☐ pH ≤2 ☐ pH ≥12.5 ☐ Solid Corrosive (WSC2)	☐ Yes	⊠ No	Unknown								
D003: Cyanide Sulfide Water Reactive Other (i.e., percodde former,	☐ Yes	⊠ No	Unknown								
D004-D043 (Identify applicable waste codes and concentrations): explosive, air reactive)	Yes ·	⊠ No	Unknown								
N/A											
<ul> <li>c) If characteristic, list any known underlying hazardous constituents (UHCs) reasonably expected to be present, and their concentrations that may be present above the LDR treatment standard (40 CFR 268.48):</li> <li>N/A</li> <li>d) List any known Land Disposal Restrictions (LDR) subcategories, if applicable (40 CFR 268.40):</li> </ul>											
e) List any applicable Washington State dangerous waste codes: (not required if federally regulated)  WT01:  Yes No Unknown  WT02: Yes No Unknown  WP01:  Yes No Unknown  WP03: List constituents and concentrations:  N/A	ixture rule fo Yes Yes Yes Yes Yes	r ignitability No No No No No	Unknown Unknown Unknown Unknown Unknown								
4. Is this material TSCA regulated for PCBs? Yes No Unknown Analysis Requested  List concentration if applicable:											
If yes, what is the source of the PCBs? (see TSCA PCB Hanford Site User Guide, DOE/RL-2001-50)											
☐ PCB Liquid Waste ☐ PCB Bulk Product Waste ☐ PCB Transformer ≥500 ppm		inknown (capacitor/b	allant) vEOO saw								
PCB Remediation Waste	· equipment (	capacitor/b	anasty Noov ppm								
5. Is this material TRU?  Yes  No  Unknown											
6 ACCURACY OF INFORMATION											
Based on my inquiry of those individuals immediately responsible for obtaining this information, that to the be-	st of my know	rledge, the l	nformation								
entered in this document is true, accurate, and complete.		8/10	7/04								
Print & Sign	Date	4//									